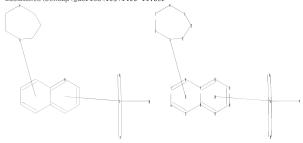
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chain nodes :
16 17 18 19
ring nodes :
1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 22 23
chain bonds :
16-17 16-18 16-19
ring bonds :
1-2 1-23 2-3 3-4 4-5 5-22 6-7 6-11 7-8 8-9 9-10 9-12 10-11 10-15
12-13 13-14 14-15 22-23
exact/norm bonds :
1-2 1-23 2-3 3-4 4-5 5-22 16-17 16-18 16-19 22-23
normalized bonds :
6-7 6-11 7-8 8-9 9-10 9-12 10-11 10-15 12-13 13-14 14-15
```

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom 11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 16:CLASS 17:CLASS 18:CLASS 19:Atom 20:Atom 21:Atom 22:Atom 23:Atom

L1 STRUCTURE UPLOADED

=> s 11

SAMPLE SEARCH INITIATED 19:35:09 FILE 'REGISTRY' SAMPLE SCREEN SEARCH COMPLETED -300 TO ITERATE

100.0% PROCESSED 300 ITERATIONS SEARCH TIME: 00.00.01

0 ANSWERS

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 4961 TO 7039
PROJECTED ANSWERS: 0 TO 0

L2 0 SEA SSS SAM L1

=> s 11 sss full

FULL SEARCH INITIATED 19:35:16 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 6203 TO ITERATE

100.0% PROCESSED 6203 ITERATIONS 3 ANSWERS

SEARCH TIME: 00.00.01

L3 3 SEA SSS FUL L1

=> file caplus

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=> s 13

1 L3

=> d 14 bib abs hitstr

- L4 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2008 ACS on STN
- AN 2003:777764 CAPLUS
- DN 139:292163
- TI Preparation of arylsulfonyl(diazacycloalkyl)quinolines for treatment of CNS disorders
- IN Ahmed, Mahmood; Johnson, Christopher Norbert; Jones, Martin C.; MacDonald, Gregor James; Moss, Stephen Frederick; Thompson, Mervyn; Wade, Charles Edward; Witty, David

PA Glaxo Group Limited, UK SO PCT Int. Appl., 48 pp. CODEN: PIXXD2

DT Patent LA English

FAN.	CNT 1			
	PATENT NO.	KIND DATE	APPLICATION NO.	DATE
PI	WO 2003080580		WO 2003-EP3197	20030325
	WO 2003080580	A3 20040205		
	W: AE, AG, AL,	AM, AT, AU, AZ,	BA, BB, BG, BR, BY, BZ,	CA, CH, CN,
			DZ, EC, EE, ES, FI, GB,	
			JP, KE, KG, KP, KR, KZ,	
			MK, MN, MW, MX, MZ, NO,	
			SG, SK, SL, TJ, TM, TN,	TR, TT, TZ,
		UZ, VC, VN, YU,		
			SL, SZ, TZ, UG, ZM, ZW,	
			BE, BG, CH, CY, CZ, DE,	
			LU, MC, NL, PT, RO, SE,	
	BF, BJ, CF,		GN, GQ, GW, ML, MR, NE,	
		A1 20031002	CA 2003-2479786	20030325
			AU 2003-219103 EP 2003-714889	
			GB, GR, IT, LI, LU, NL,	
			CY, AL, TR, BG, CZ, EE,	
			BR 2003-8696	
			CN 2003-811644	
		T 20051020	JP 2003-578335	
	JP 2005531518 TW 268928	B 20061221	TW 2003-92106558	
	RU 2309154	C2 20071027	RU 2004-131641	20030325
	NZ 535239		NZ 2003-535239	20030325
	ZA 2004007320		ZA 2004-7320	20040912
	IN 2004DN02703	A 20070302	IN 2004-DN2703	20040914
	MX 2004PA09318	A 20050125	MX 2004-PA9318	20040924
	US 20050124628	A1 20050609	US 2004-509078 NO 2004-4588	20040927
	NO 2004004588	A 20041025	NO 2004-4588	20041025
PRAI	GB 2002-7289	A 20020327		
	GB 2002-25678			
		W 20030325		
os	MARPAT 139:292163			
GI				

AB Title compds. I [R1, R2 = H, alky1; R1R2, R22 = (CH2)2-4; R3-R5 = H, halogen, CN, CF3, OCF3, alky1, alkoxy, alkanoy1, (un) substituted COMH2; A = (un)substituted cary1; m = 1-4; n = 1-3, p = 1, 2] were prepared for use as H16 receptor antagoniets in treatment of CNS disorders. Thus, 8-iodo-3-phenylsulfonylquinoline was prepared from 8-nitroquinoline and was treated with 1-tert.-butoxycarbonylpiperazine, followed by deblocking, to give 3-phenylsulfonyl-8-piperazineoquinoline.

607743-12-4P RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of arylsulfonyl(diazacycloalkyl)quinolines for treatment of CNS disorders)

RN 607743-12-4 CAPLUS

CN 1H-1,4-Diazepine-1-carboxylic acid, hexahydro-4-[3-(phenylsulfonyl)-8-quinolinyl]-, 1,1-dimethylethyl ester (CA INDEX NAME)

IT 607742-67-6P 607743-57-7P

RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of arylsulfonyl(diazacycloalkyl)quinolines for treatment of CNS disorders)

RN 607742-67-6 CAPLUS

CN Quinoline, 8-(hexahydro-1H-1,4-diazepin-1-y1)-3-(phenylsulfonyl)-, hydrochloride (1:1) (CA INDEX NAME)

● HC1

RN 607743-57-7 CAPLUS CN Quinoline, 8-(hexahydro-1H-1,4-diazepin-1-yl)-3-(phenylsulfonyl)- (CA INDEX NAME)

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10/571405

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